Format for Detailed Project Information

1. **Project Title :**
2. **Introduction:** (Introduce the subject. Provide full background information. Cite literatures that are specific to the topic of the research proposal. Information should be completed to prove that the research proposal is based on a sound scientific footing.)
3. **Rationale:** (Describe the relevance of the proposed study to national health priorities and relationship of the objectives to existing scientific knowledge on the subject. Cite relevant literature and refer to related studies done in our country or elsewhere.)
4. **Hypothesis or research question:**
5. **Objectives:** (List the general and specific objectives of the proposed study and state clearly the question that is being posed or the hypothesis being tested.)
6. **Methodology:** Describe the design of the study and methodology in sufficient detail to enable assessment of how they will contribute towards achievement of the stated objectives and to permit proper appraisal of the budget. Plan for data analysis to be included if relevant and important. This section should contain details of the research methods. Enough detail should be given to evaluate whether the methods are already tested and feasible. The following scheme is suggested:
   Factors in study (variables), Study Population, Sampling, Statistical basis of the sample size, Procedures, Methods of Data Collection, Pretesting, Data Interpretation, Statistical (Correlation, Significance Test, Coefficient of Variation, Evaluation, Methods, wherever applicable).
7. **Utilization of Results:** (Describe in brief how you perceive that the results from this may contribute to health development of the Country.)
8. **Facilities :** (Resources, equipment, chemicals, subjects (human, animal) etc. required for the study):
9. **Additional Facilities Required** :
10. **Flow Chart:** (Describe sequence of tasks within time frame)
11. **Budget of research-**
12. **Ethical Implications:** (Think very carefully about possible ethical implications and put views. Consult BMRC's Guidelines for Ethical Review of Projects involving Human Subjects).

13. Consent Form – Both English and Bengali

14. Approval / Forwarding of the Head of Department / Institute / IRB

15. **Mutual of understanding (MOU) documents –** From DGDA, Drug producing agencies / Institutional administrative authority wherever needed.
16. **References:** Vancouver / Harvard style to be followed. e.g.- Can Med Assoc J 1995; 152(9):1459-1465.